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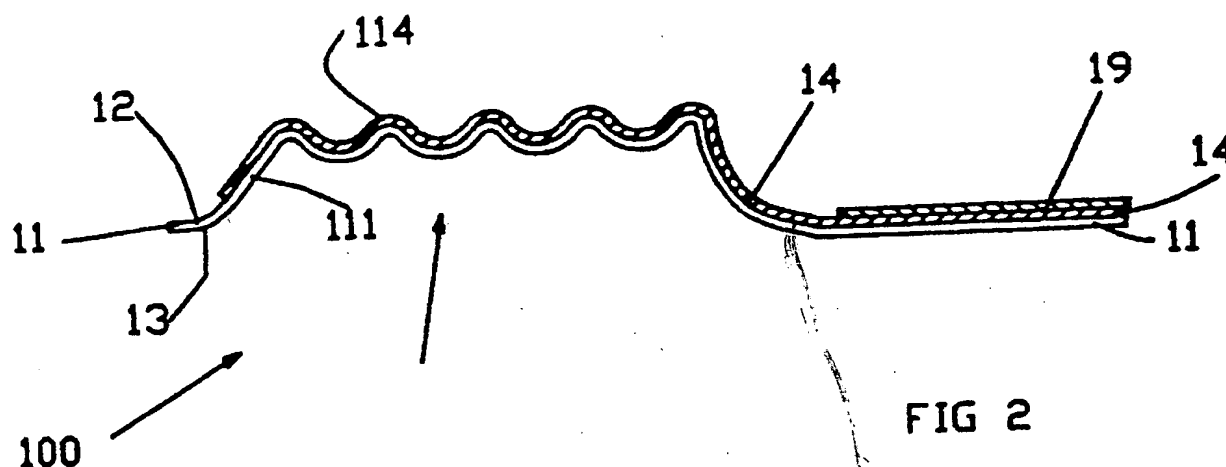
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(54) Profiled biosignal electrode device

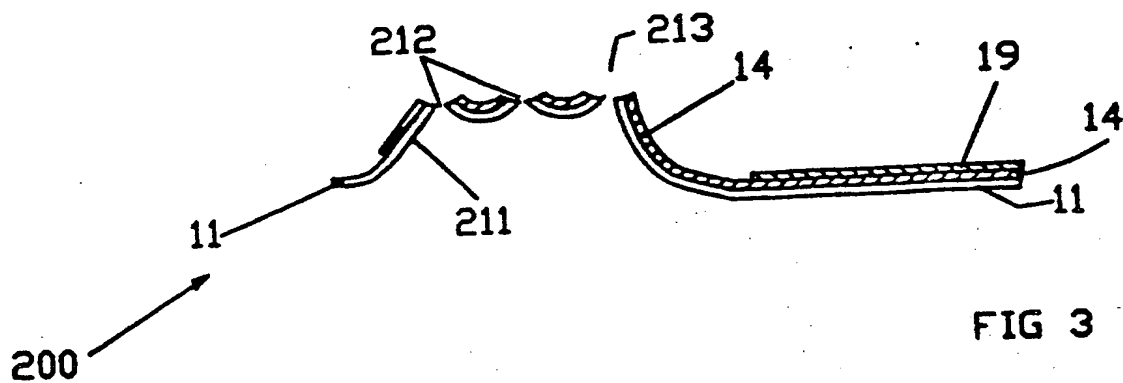
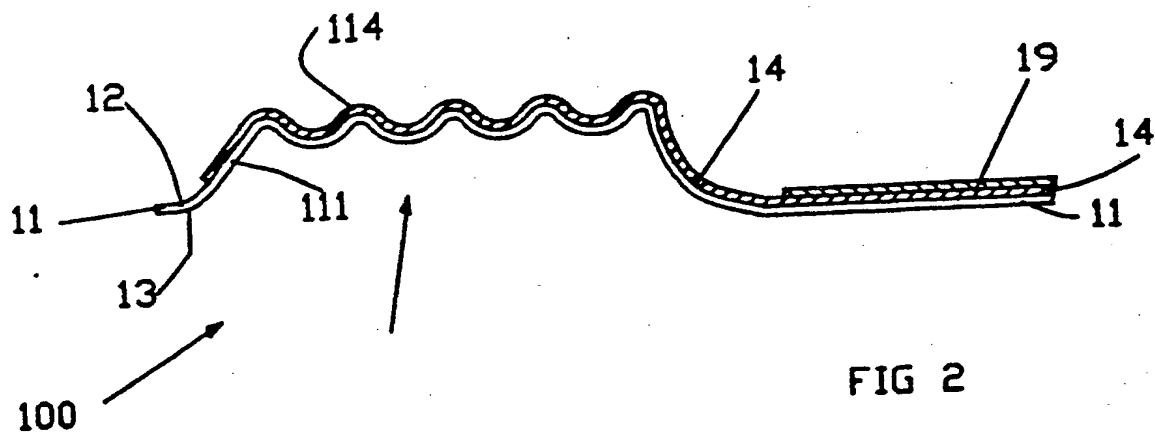
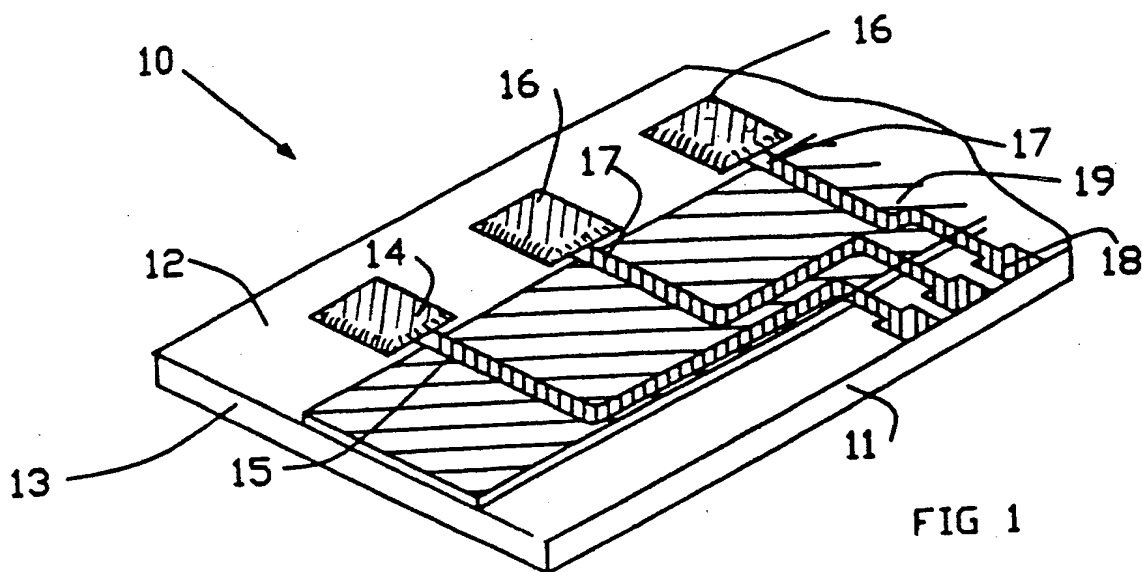
(57) A biosignal electrode device comprises a flexible electrically insulating substrate 11 having deposited thereon an electrically conductive layer 14 forming an electrode sensor 114 and a lead for the sensor. The portion of the substrate bearing the sensor 114 is formed in relief to improve electrical contact with the skin. Recesses in the face of the sensor may accommodate conductive gel or the relief portions may be punctured to form fine points. A plurality of electrodes may be formed on the same flexible substrate.

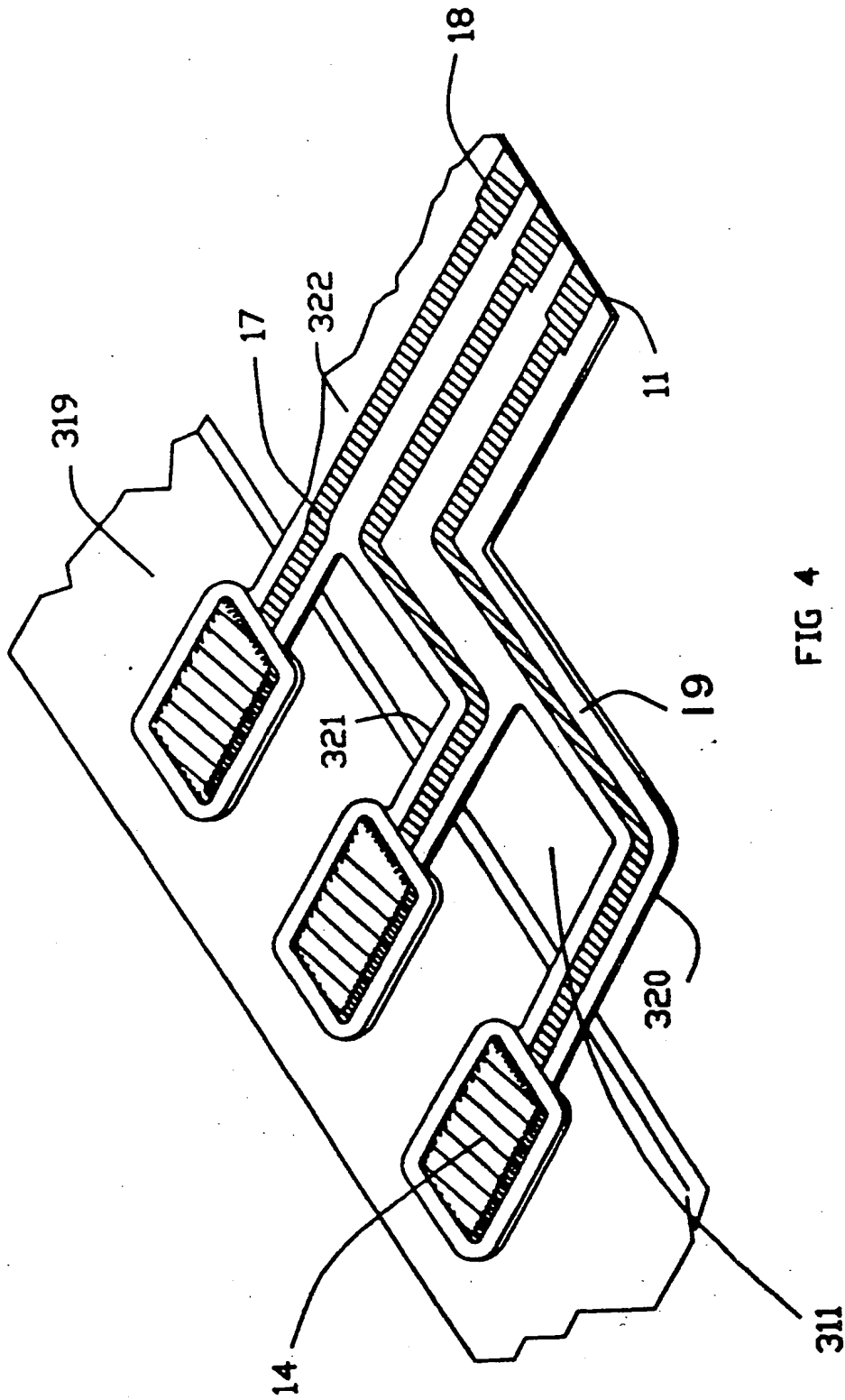


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At least one drawing originally filed was informal and the print reproduced here is taken from a later filed formal copy.

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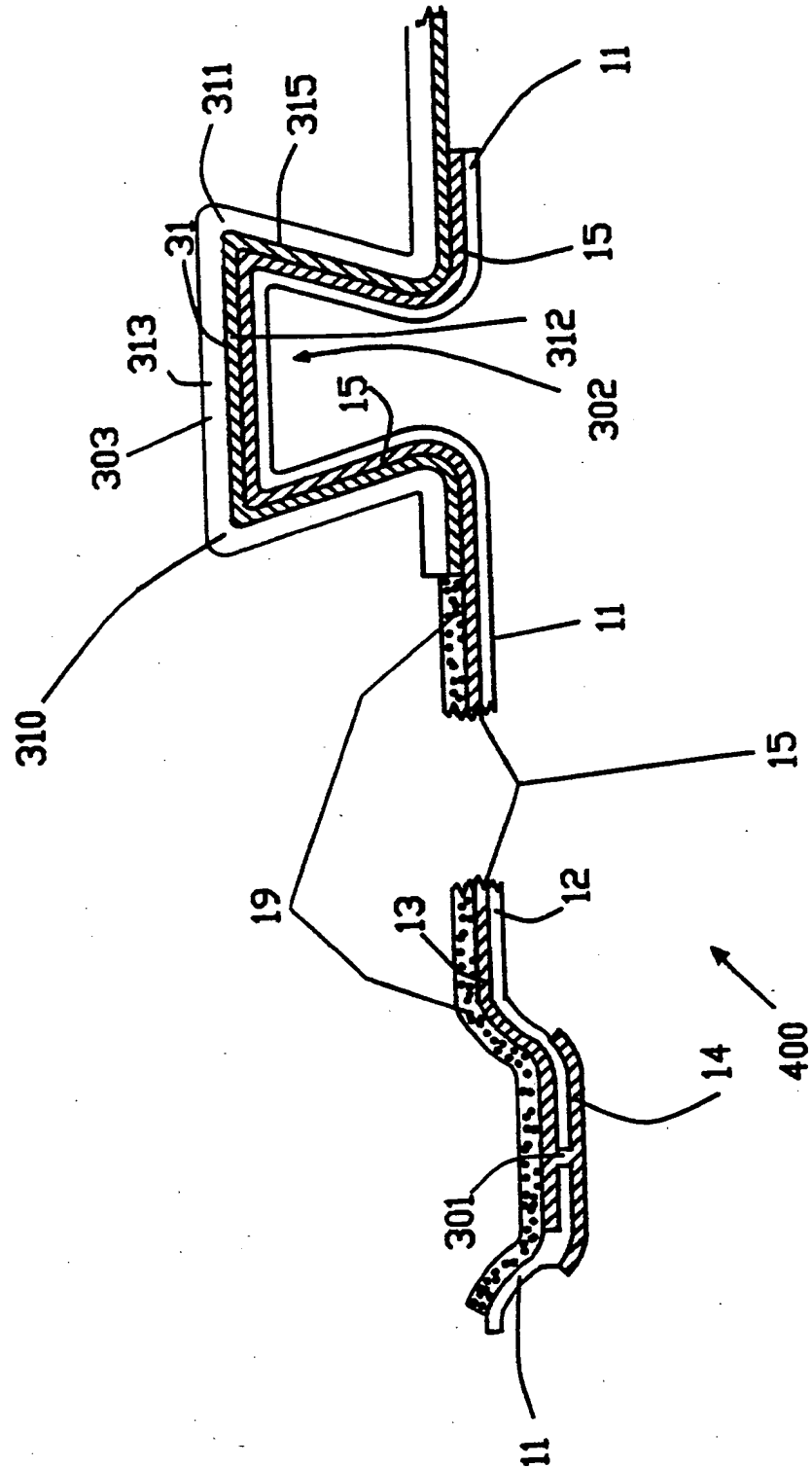
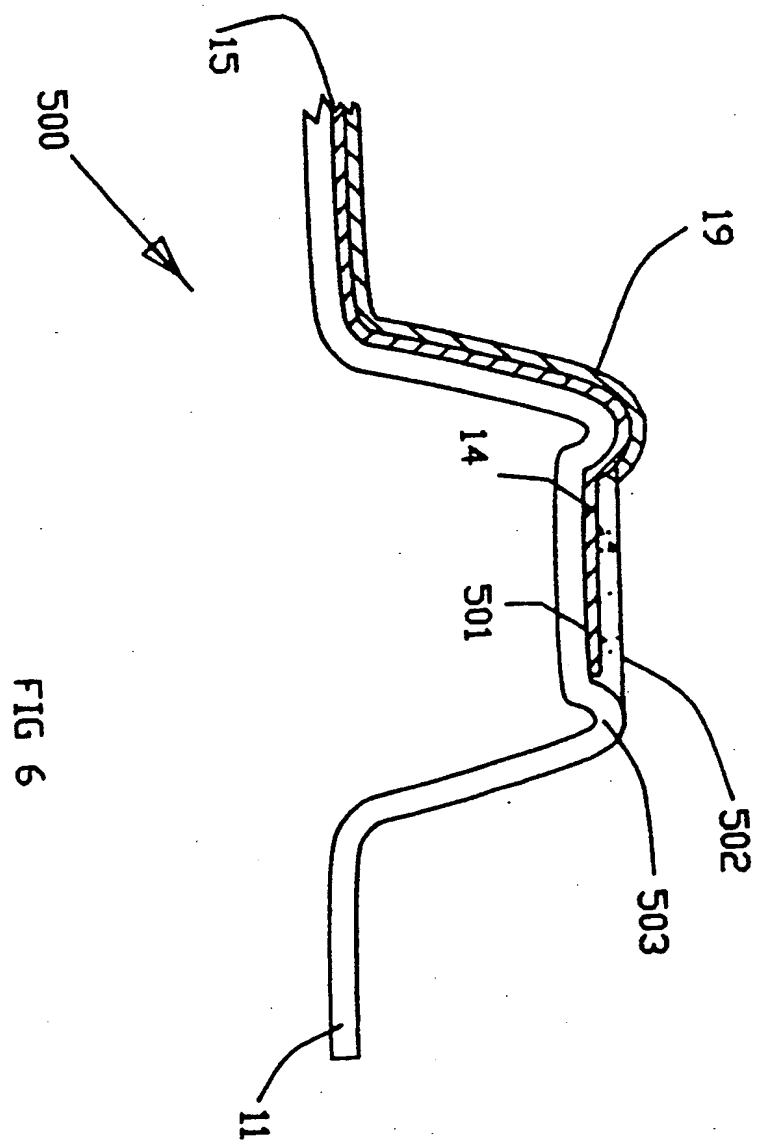


FIG 5



- 1 -

PROFILED BIOSIGNAL ELECTRODE DEVICE

This invention relates to a profiled biosignal electrode device.

5 The skin's outermost layer, the stratum corneum, is composed of dead cells. It is relatively nonconductive and presents a high impedance to the transmission of bioelectric signals. Stratum corneum, due to its dielectric properties and thinness, permits capacitive coupling between a metal electrode placed on the skin surface and the underlying
10 conductive tissues. Some ions manage to traverse the stratum corneum, and the flow of this ionic current can be represented electrically by a large resistance shunting the skin's capacitance. If the electrode-skin impedance is large relative to the input impedance of the amplifier,
15 biosignal attenuation occurs. The presence of a reactive component in the electrode-skin impedance can also create filtering and, hence, distortion of the signal.

20 Another problem source is 50/60Hz electric-field coupling between neighbouring powerlines and the patient's monitoring cables. This interference is due to electrode-skin impedance mismatch, which leads to a differential interference signal at the amplifier's input.

Variance in ionic concentration across the epidermis causes a potential difference. This potential has a typical value of 30mV, with the skin surface normally negative relative to the inside of the body. This potential varies
5 from site to site, patient to patient, and depends on gel composition which is used in conjunction with the electrode and skin condition. Slight differences in electrode materials and skin-site parameters cause electrode-skin potentials to be mismatched. The difference between
10 potentials will be amplified along with the desired biosignal. Fluctuations in the magnitudes of potentials and their difference can introduce an undesired frequency component into the desired signal, making accurate diagnosis difficult. The dependence of the electrode-skin potential
15 on thickness of the epidermal layer is important in many biosignal recording applications. If that thickness is changed by stretching or pressing on the skin, the potential can vary by up to 10mV. This is the major source of motion artefact (fluctuations in baseline trace with patient move-
20 ment) and a function of the degree of skin deformation.

Many current electrodes incorporate electrolytic gels which serve to ensure an optimal electrical contact between the electrode sensor and the patient's skin and which also
25 serve to decrease the high epidermal impedance.

The use of electrolytic gels not only decreases the skin's impedance, thus avoiding associated problems, but also has been observed to reduce motion artefact levels.
30

The potential disadvantages of using gelled electrodes reside in the utilization of the electrolyte gel which requires that such gel be extraneously supplied during application. Usually, the gel is packed in a tube which must be

squeezed for dispensing a charge thereof. Sometimes, a charge of gel is packaged separately to that of the electrode in an hermetically sealed container. In the former case, the necessity to provide extraneous gel during application is an obvious inconvenience, especially in the home-based environment under emergency conditions. Following use, the electrode (if it is reusable) and the application site must be cleaned.

In the latter case, once the package is opened to atmospheric conditions the gel will rapidly dry out. Even when properly packaged, the gelled electrode will have a relatively short shelf life. For home-based monitoring applications, for example, a sufficient supply of such pregelled disposable packages must be provided and must be periodically renewed.

A simple method of avoiding the above problem in, for example, the home-based monitoring environment, is to use dry metal plate electrode or electrodes attached, for example, to a limb of the patient by means of elasticated straps. Although no gel is used, the accumulation of perspiration under the metallic plate will eventually moisten the skin covered by the 'dry' electrode. The initial impedance is however much higher than that for a gelled electrode and the impedance decreases relatively slowly with time. This need not be a major problem, however, as amplifiers now exist with input impedances and common-mode rejection ratios sufficiently high to cope with such large electrode-skin impedances.

The advantage of such electrolyte-free electrodes are numerous. The electrodes can be applied rapidly and with ease to a patient's limb as they do not require skin or

electrode preparation prior to or after electrode application. The electrodes are reusable, pose no storage problems and there is little corrosion of the metal plate. Such electrodes have in the past proved optimal, for
5 example, for low cost, rapid, mass screening of electrocardiograms.

Dry metal plate electrodes are however susceptible to 50/60Hz interference problems due to the mismatching of
10 their high electrode-skin impedances. They do not perform well on hairy subjects due to poor electrical contact. Even when firmly strapped to a part of the body, a rigid, metal plate electrode is prone to generate motion artefacts when the underlying 'dry' skin is stretched or moved or when the
15 electrode is pressed. The relatively rigid plate electrode is only suitable as a limb electrode as it cannot easily be attached to a patient's chest, for example, and is incapable of accommodating body contours.

20 It is an object of the present invention to overcome these problems.

The invention, therefore, provides a biosignal electrode device comprising a flexible electrically insulating
25 substrate having deposited thereon an electrically conductive layer forming an electrode sensor and a lead for the sensor, the portion of the substrate bearing the sensor being formed in relief.

30 Preferably, the surface of the sensor element is rough, textured, rippled, punctured or the like, all of which are hereinafter embraced by the word "rough".

The biosignal electrode device according to the invention may be manufactured by several well-known techniques including embossing, debossing or hydroform embossing. In the embossing or debossing processes, matching male and female die sets are used to form the material. As it is the male die projections which apply the initial pressure to the unsupported substrate, they give rise to unequal strains in the substrate which can distort the form shape. Problems with thermal expansion may be encountered with thermal forming processes and tooling temperature can cause damage to the inking systems. Accordingly, the preferred manufacturing technique is by way of hydroform embossing.

In hydroform embossing, one half of the conventional die set is eliminated and replaced with a hydraulic fluid cell assembly which has a urethane diaphragm face. Hydroform embossing therefore uses a single sided tool with the fluid cell assembly serving as the active compressive "counter-die". With hydroform embossing, hydrostatically equalized holding forces are applied to the entire die surface area and substrate before forming into the die cavity occurs. The substrate may therefore be stretched-formed into the cavity rather than gathering around male die projections and being forced or pinched into the female tool, with resulting damage to the substrate. A counter die in the form of an elastomer diaphragm will not mar the polymer substrate.

A further advantage of hydroform embossing is that the one sided tool can be created directly from photo-artworks and thus permits the accurate registration from the embossing artwork to the tool. The single sided tool also allows substrate to be accurately pin-registered to the die plate.

With hydroform embossing, many unique embossing applications are possible which cannot be achieved by conventional techniques such as the forming of angular bends, the deep drawing of shapes and the creation of negative or zero draft angles on the formed part.

In order to provide the roughness in the surface of the sensor element the substrate and the conductive layer must be suitably malleable. Further, the coefficient of thermal expansion of the conductive layer should match that of the flexible substrate. The flexibility of the materials should be such as to minimise cracking, peeling etc.

The conductive layer may be provided by means of an electrically conductive ink which also has the property of "self protecting" against corrosion and residual growth on the surface.

A suitable thermoplastic material loaded, for example, with silver and silver-chloride particles may be used as the conductive layer. Appropriate choice of particle size, thermoplastic binder, deposition and curing parameters leads to an optimal conductive layer surface ensuring good electrical contact with the skin.

Moulded surface "ripples" or punctured "points" in the sensor further optimise contact between the sensor and the patient's skin. Due to frictional forces, the ripples or points minimise the movement of the electrode relative to skin and thus ensure more stable biosignal recording. The ripples or points increase the contact area and, when pressed against the skin, penetrate the epidermal layer sufficiently to decrease significantly the electrode-skin

potential fluctuations (motion artefact) without
traumatising the skin.

5 The substrate surrounding the lead may be cut or
otherwise removed in order to leave a finger of substrate on
which is located the lead. Such a finger enables relative
movement of the sensor and thus accommodates changes in the
monitored body segment's dimensions and form due to
breathing, patient movement etc.

10 Multiple electrode devices, referred to herein as
"Off-shoot" electrodes may be connected to a common
electrode harness. The harness may be formed with multiple
female connectors which may be made by hydroform embossing
15 in areas of the harness substrate coated with a conductive
layer. The off-shoot electrodes may then be connected by
means of male connectors, also manufactured by hydroform
embossing, to respective ones of the female connectors on
the harness. The conductive layer forming the connecting
20 lead on, for example, the obverse face of the "off-shoot"
electrode is through-hole-plated to the reverse face of the
substrate in order to make connection with the electrode
sensor. This method of connection enables the off-shoot
electrode to pivot about the connector enabling the
25 electrode device to accommodate anatomical inter-human
variations. Several sections incorporating male and female
connectors at each end may be connected in series in order
to produce articulated off-shoot electrodes which have a
high degree of manoeuvrability. Good electrical contact can
30 be ensured by: (i) providing a tight male-female connector
fit: (ii) by roughening the connecting surfaces of the male
and female connectors during the forming process; and (iii)

by depositing conductive layers at the connector sites which have rough surfaces due, for example, to the particle size of the filler.

5 The conductive layer may be printed or otherwise deposited onto the substrate as required for a given application. Where there are multiple sensors the sensors may be backed by a band of, for example, suitable foam in order to
10 (i) keep the sensor elements in their correct relative positions when the surrounding substrate is removed (ii) make the electrode device more manageable and avoid lead entanglement (iii) help distribute any applied pressure evenly over the electrodes and (iv) thus help minimise motion artefact. The backing layer should preferably be
15 lightweight, flexible and stretchable to ensure that the harness is comfortable, that it can conform readily with body contours and that it can accommodate changes in the body segment's dimensions due to breathing or patient movement.

20

 The electrode device according to the invention may be applied manually by the subject or clinician to the part of the body under study in applications which only require the short term, "diagnostic" observation of the detected
25 biosignal(s) and without the need for a gel layer. For longer term monitoring applications an electrode harness comprising a plurality of electrode sensor elements may be held in place on the required body part by means of elasticated straps, adhesive tape, suction cups etc. and
30 again without the need for a gel layer.

 The electrode device according to the invention when in the form of a harness may be gelled, preferably with hydrogel pads, for certain biomedical applications. "Solid",

conductive, adhesive hydrogel pads can be coated onto the
profiled printed sensors, giving rise to a simple, easy-to-
manufacture electrode structure. The use of such hydrogel
dispense with the need for the gel-retaining ring,
5 gel-impregnated sponge and sealing cap used in current 'wet'
gel electrodes.

The use of adhesive hydrogel on the electrode device
according to the invention is advantageous in applications
10 where, for example, large numbers of electrodes must be
applied to the body. Body surface cardiac mapping is one
such application. The number and distribution of the
electrodes is such that it would be difficult to hold dry
electrodes sufficiently well, manually or by means of
15 straps, to ensure good electrical contact.

The roughened surface of the electrode sensor element
ensures firm electrical contact between the sensor and the
gel pad as well as increasing the contact area and thus
20 decreasing the sensor-gel interface impedance. That part of
the electrode device having the electrode sensor element
thereon may be contoured during the embossing technique in
such a way as to form a recess around the sensor element
which is surrounded by a lip and into which recess gel may
25 be loaded. The recess would enable gel, while in a fluid
state, to be poured onto the profiled sensor and cured *in situ*.
Following the application of the gel into the
recesses, a removable backing strip may be set in place so
as to retain the integrity of the gel prior to use. The
30 electrode device according to the invention when in the form
of a harness enables a plurality of sensor elements to
conform optimally with the body's contours.

The invention will be understood in greater detail from

the following description of preferred embodiments thereof given by way of example only and with reference to the accompanying drawings in which:-

5 FIGURE 1 is a perspective view of a first embodiment of a biosignal electrode according to the invention;

FIGURE 2 is a cross-sectional view of a second embodiment of a biosignal electrode according to the
10 invention;

FIGURE 3 is a cross-sectional view of a third embodiment of a biosignal electrode according to the invention;

15 FIGURE 4 is a perspective view of a fourth embodiment of a biosignal electrode according to the invention; and

FIGURE 5 is a cross-sectional view of a fifth embodiment of a biosignal electrode according to the
20 invention; and

FIGURE 6 is a cross-sectional view of a sixth embodiment of a biosignal electrode according to the invention.

25 Referring now to the drawings, and in particular to Figure 1 thereof, there is shown a first embodiment of an electrode device 10 according to the invention which comprises a substrate 11 having an obverse face 12 and a reverse face 13. Onto the obverse face 12 is deposited an
30 electrode sensor 14 and an associated lead 15. Additional sensors 16 and respective associated leads 17 may also be deposited on to the obverse face 12 or the reverse face 13 if desired. Each of the leads 15, 17 terminates in a respective connector end 18. The sensors 14, 16; leads 15, 17; and connector ends 18 may be deposited by well-known

techniques including screenprinting of a suitable electrically conductive ink. A dielectric layer 19 is deposited over the leads 15, 17 in order to insulate them from a patient on which the device 10 will be used. It will be appreciated that the sensors 14, 16 and the connector ends 18 are allowed to remain exposed. The parts of the substrate 11 having the sensors 14, 16 thereon are in relief relative to the remainder of the substrate 11. To provide for such relief, the substrate may be subject to well-known techniques such as profiling, embossing, forming, moulding or casting. A preferred technique is that of hydroform embossing as discussed earlier in the present specification.

In Figure 2 of the drawings, there is shown a second embodiment of an electrode device 100 according to the invention which is substantially similar to the electrode device 10 save as follows. That part 111 of the substrate 11 which is in relief is rippled or indented and consequently the electrode sensors 14, 16 are also rippled or indented at 114 thereby giving rise to a rough or textured surface to the electrode sensors 14, 16.

In Figure 3 of the drawings, there is shown a third embodiment of an electrode device 200 according to the invention which is substantially similar to the electrode device 10 except as follows. The part 211 of the substrate 11 which is in relief is punctured with a plurality of relatively small openings 212 which give rise to a plurality of fine points 213 on the electrode sensors 14, 16.

30

The use of a thin, flexible substrate 11, and dielectric layer 19 layers gives rise to a flexible overall electrode device according to the invention capable of conforming with the body's contours. The protrusion of the electrode sensor

14, 16, further optimises the contact between the electrode sensors 14,16 and the subject's skin when the device 10 is applied to the skin. With the electrode sensors 14,16 in relief, there is provided the additional advantageous feature of absorbing the effects of variations in pressure on the device according to the invention when in use and thus help minimise motion artefacts which are generally a major problem with dry electrode structures.

10 In Figure 4 of the drawings, there is shown a fourth embodiment of a device 300 according to the invention which is substantially similar to the device 10 of Figure 1 of the drawings and which may have the additional features as described with respect to devices 100 or 200 except as follows.

15 The part of the substrate 311 which surrounds the leads 17 may be cut or otherwise removed in order to leave fingers 320, 321, 322 of substrate on which are located the leads 17. The provision of the fingers 320, 321, 322 enables movement of the sensor elements 14, 16 relative to the connector ends 18 and thus accommodates movement of the subject due, for example, to breathing etc.

25 The sensors 14, 16 and associated leads 15 may be printed or otherwise deposited onto the substrate 11 at the required relative distances and orientations for a given application. Where possible, the sensors 14, 16 may be backed by a band 319 of, for example, suitable foam in order to (i) keep the sensors 14, 16 in their correct relative positions when the surrounding substrate 11 is removed (ii) make the electrode device 300 more manageable and avoid lead entanglement (iii) help distribute any applied pressure evenly over the sensors 14, 16 and (iv) thus help minimise

motion artefact. The band 319 should preferably be
lightweight, flexible and stretchable to ensure comfort;
conformity with body contours; and to enable changes in the
body segment's dimensions due to breathing or patient move-
5 ment to be accomodated.

In Figure 5 of the drawings which relates to an
"off-shoot" electrode, there is shown a fifth embodiment 400
of an electrode device according to the invention which is
10 substantially similar to the device 10 of Figure 1 of the
drawings except as follows. First, it comprises only a
single sensor 14 and lead 15. Second, the lead 15 is absent
from the obverse face 12 being, instead, deposited on the
reverse face 13. The lead 15 is in electrical communication
15 with the sensor 14 having regard to the provision of a hole
301 in the substrate 11 which enables the electrically
conductive ink to flow through the hole 301 thereby
providing electrical communication between the sensor 14 and
the lead 15. Further, the end of the substrate 11 which is
20 remote from the end having the sensor 14 thereon is formed
into a u-shape in a direction opposite to that of the
substrate 11 on which the sensor 14 is deposited to provide
a male connector 302. The reverse face 13 of the substrate
11 having the male connector 302 thereon also has a
25 continuation of the lead 15 thereon.

A harness 310 is provided which essentially comprises a
substrate 311 also having an obverse face 312 and a reverse
face 313. The harness has a plurality of shapes in the form
30 of female connectors 303 which are engageable with
respective male connectors 302.

The obverse face 312 of the harness 310 has a lead 315
deposited thereon in a manner similar to that of the lead
15. The lead 315 extends into the female connector

303 so that when the connectors 302 and 303 are mating as shown in Figure 5, there is electrical connection between the leads 15 and 315. The lead 315 may terminate on the harness 310 in a manner similar to the connector end 18. It
5 will be appreciated that the harness 310 will have a plurality of leads 15 in electrical isolation from each other to enable a plurality of off-shoot electrode devices to be connected to the harness so that the harness 310 not only provides a means for enabling a plurality of electrode
10 devices to be attached to the body of a patient or subject but also to provide an independent electrical lead from each electrode device the ends of which leads may be individually electrically connected to a suitable monitoring device.

15 In Figure 6 of the drawings, there is shown a fifth embodiment of an electrode device 500 according to the invention which is essentially similar to the device shown in Figure 1 of the drawings except as follows. That part of the substrate 11 having the sensor 14 thereon which is in
20 relief is provided with a well 501 having a lip 503 and in which the sensor element 14 is located. Following the manufacture of the electrode device, a suitable electrolytic gel 502 is poured into the well 501 which, depending on the type of gel used, cures to a viscous consistency. There is
25 then applied to the well 501 or more preferably a plurality of wells a backing strip which is removable prior to the use of the device.

30 The invention is not limited by or to the specific embodiment described which can undergo considerable variation without departing from the scope of the invention.

CLAIMS:

1. A biosignal electrode device comprising a flexible electrically insulating substrate having deposited thereon an electrically conductive layer forming an electrode sensor and a lead for the sensor, the portion of the substrate bearing the sensor being formed in relief.
2. A device as claimed in claim 1, wherein the surface of the sensor is roughened.
3. A device as claimed in claim 1 or 2, wherein the lead is deposited on the opposite surface of the substrate to the sensor and is connected to the sensor through an aperture in the substrate.
4. A device as claimed in claim 1, 2 or 3, wherein the portion of the substrate which is in relief and bears the sensor is slightly recessed to form a well, and the well contains an electrolytic gel.
5. A device as claimed in claim 1, 2, 3 or 4, wherein there is a plurality of sensors and associated leads on the substrate.
6. A device as claimed in claim 5, wherein regions of the substrate between the sensors and leads are removed to provide fingers of substrate each bearing a sensor and its associated lead.
7. A device as claimed in claim 6, wherein the sensors are backed by a flexible and stretchable band of insulating material.

8. A device as claimed in claim 1, 2, 3 or 4,
wherein a portion of the substrate remote from the
sensor is formed into one part of a male/female
connector, the connector part having an electrically
5 conductive layer thereon which is electrically
connected to the lead.

9. A biosignal electrode assembly comprising a
harness in the form of an electrically insulating
10 substrate having a plurality of leads deposited thereon
each terminating in one part of a male/female
connector, and a plurality of biosignal electrode
devices as claimed in claim 8 each connectable by its
connector part to a respective connector part on the
15 harness.

10. A biosignal electrode device substantially as
hereinbefore described with reference to, and as illus-
trated in, the accompanying drawings.

Patents Act 1977
Examiner's report to the Comptroller under Section 17
(The Search report)

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17-

Relevant Technical Fields

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(ii) Int Cl (Ed.5) A61B 5/0408, 5/0416, 5/0478, 5/0492, 5/05;
A61N 1/02, 1/04, 1/06

Databases (see below)

(i) UK Patent Office collections of GB, EP, WO and US patent specifications.

(ii) ONLINE DATABASE: WPI

Search Examiner
R S CLARK

Date of completion of Search
23 JUNE 1994

Documents considered relevant
following a search in respect of
Claims :-
1,2

Categories of documents

- X:** Document indicating lack of novelty or of inventive step. **P:** Document published on or after the declared priority date but before the filing date of the present application.
- Y:** Document indicating lack of inventive step if combined with one or more other documents of the same category. **E:** Patent document published on or after, but with priority date earlier than, the filing date of the present application.
- A:** Document indicating technological background and/or state of the art. **&:** Member of the same patent family; corresponding document.

Category	Identity of document and relevant passages		Relevant to claim(s)
X	GB 2160107 A	(ASTRA-TECH) page 2 lines 10-15	1,2
X	GB 1575364	(RASMUSSEN) whole document	1,4,5,6,8
X	GB 1128329	(NRDC) page 2 lines 92-129	1,2
X	EP 0014341 A1	(QUINTON) pages 5-6	1,2
X	US 3989036	(SASAMORI) Figure 4	1,2
X	US 3741219	(SESSIONS) whole document	1,2

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